

## CLAIMS

What is claimed is:

1. A method for preventing or for treating chronic pain comprising administering to a patient in need of treatment an effective amount of a synergistic combination of a NK<sub>1</sub> receptor antagonist and a GABA analog.
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2. A method of Claim 1 wherein the ratio of the GABA analog relative to the NK<sub>1</sub> receptor antagonist is from 50:1 to 1:1 expressed as parts by weight.
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3. A method of Claim 1 wherein the ratio of the GABA analog relative to the NK<sub>1</sub> receptor antagonist is 20:1 expressed as parts by weight.
4. A method according to Claim 1 wherein the NK<sub>1</sub> receptor antagonist is [2-(1*H*-indol-3-yl)-1-methyl-1-(1-phenyl-ethylcarbamoyl)-ethyl]-carbamic acid benzofuran-2-ylmethyl ester [R-(R\*,S\*)].
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5. A method according to Claim 1 wherein the GABA analog is gabapentin.
6. A method according to Claim 1 wherein the GABA analog is pregabalin.
7. A method according to Claim 1 employing [2-(1*H*-indol-3-yl)-1-methyl-1-(1-phenyl-ethylcarbamoyl)-ethyl]-carbamic acid benzofuran-2-ylmethyl ester [R-(R\*,S\*)] and gabapentin.
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8. A method according to Claim 1 employing [2-(1*H*-indol-3-yl)-1-methyl-1-(1-phenyl-ethylcarbamoyl)-ethyl]-carbamic acid benzofuran-2-ylmethyl ester [R-(R\*,S\*)] and pregabalin.
9. A method according to Claim 1 wherein the condition treated is selected from causalgia, neuropathic pain, diabetic neuropathy, post-surgery or

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traumatic neuropathy, postherpetic neuralgia, peripheral neuropathy, entrapment neuropathy, phantom limb and stump pain, neuropathy caused by alcohol abuse, HIV infection, multiple sclerosis, hypothyroidism or anticancer chemotherapy

5 10. A pharmaceutical composition comprising synergistic effective amounts of a NK<sub>1</sub> receptor antagonist and a GABA analog.

11. A composition of Claim 10 wherein the ratio of the GABA analog relative to the NK<sub>1</sub> receptor antagonist is from 50:1 to 1:1 expressed as parts by weight.

10 12. A composition of Claim 10 wherein the ratio of the GABA analog relative to the NK<sub>1</sub> receptor antagonist is 20:1 expressed as parts by weight.

13. A composition of Claim 10 wherein the NK<sub>1</sub> receptor antagonist is [2-(1*H*-indol-3-yl)-1-methyl-1-(1-phenyl-ethylcarbamoyl)-ethyl]-carbamic acid benzofuran-2-ylmethyl ester [R-(R\*,S\*)].

15 14. A composition of Claim 10 wherein the GABA analog is gabapentin.

15. A composition of Claim 10 wherein the GABA analog is pregabalin.

16. A composition of Claim 10 employing [2-(1*H*-indol-3-yl)-1-methyl-1-(1-phenyl-ethylcarbamoyl)-ethyl]-carbamic acid benzofuran-2-ylmethyl ester [R-(R\*,S\*)] and gabapentin.

20 17. A composition of Claim 1 employing [2-(1*H*-indol-3-yl)-1-methyl-1-(1-phenyl-ethylcarbamoyl)-ethyl]-carbamic acid benzofuran-2-ylmethyl ester [R-(R\*,S\*)] and pregabalin.

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18. The use of a composition comprising synergistic effective amounts of a NK<sub>1</sub> receptor antagonist and a GABA analog, or pharmaceutically acceptable salts thereof, for the preparation of a medicament useful for preventing or treating chronic pain.